CLAIMS

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- 1. An immunogenic composition comprising:
 - (a) an immunostimulating amount of a Neisseria antigen; and
 - (b) an immunostimulating amount of an adjuvant composition comprising an oligonucleotide comprising at least one CG motif.
- 2. The composition of claim 1, wherein said Neisseria antigen is selected from the group consisting of a protein, protein-polysaccharide, protein-lipopolysaccharide, polysaccharide, and lipopolysaccharide.
- 3. The composition of any preceding claim, wherein said Neisseria antigen is from *Neisseria meningitidis* or *Neisseria gonorrhoeae*.
 - 4. The composition of claim 3 wherein said Neisseria antigen is a *Neisseria meningitidis* serogroup B peptide.
 - 5. The composition of claim 4 wherein said peptide comprises SEQ ID NO:31.
- 6. The composition of any preceding claim, wherein component (b) further comprises a second adjuvant.
 - 7. The composition of claim 6, wherein said second adjuvant comprises an oil droplet emulsion.
 - 8. The composition of claim 7, wherein said oil droplet emulsion comprises a metabolizable oil and an emulsifying agent.
- 9. The composition of claim 8, wherein said oil and said emulsifying agent are present in the form of an oil-in-water emulsion having oil droplets substantially all of which are less than 1 micron in diameter and wherein said composition exists in the absence of any polyoxypropylene-polyoxyethylene block copolymer.
- 10. The composition of claim 9, wherein said oil is an animal oil, an unsaturated hydrocarbon,a vegetable oil, or a terpenoid.
 - 11. The composition of claim 10 wherein said terpenoid is squalene.
 - 12. The composition of any one of claims 8 to 11, wherein said composition comprises 0.5 to 20% by volume of said oil in an aqueous medium.

- 13. The composition of any one of claims 8 to 12, wherein said emulsifying agent comprises a non-ionic detergent or a polyoxyethylene sorbitan mono-, di-, or triester or a sorbitan mono-, di-, or triether.
- 14. The composition of any one of claims 8 to 13, wherein said composition comprises 0.01 to 0.5 % by weight of said emulsifying agent.

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- 15. The composition of any preceding claim, further comprising a separate immunostimulating agent.
- 16. The composition of claim 15 wherein said immunostimulating agent is selected from the group consisting of a bacterial cell wall component and muramyl peptide.
- 17. The composition of any one of claims 6 to 17, wherein said second adjuvant comprises alum, incomplete Freund's adjuvant (IFA), or complete Freund's adjuvant (CFA).
 - 18. The composition of any preceding claim, wherein said oligonucleotide comprises at least one phosphorothioate bond.
- 19. The composition of any preceding claim, wherein said oligonucleotide comprises a nucleotide sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, and SEQ ID NO:27.
 - 20. The composition of any preceding claim, wherein said oligonucleotide comprises a CG motif flanked by two purines immediately 5' to said motif and two pyrimidines immediately 3' to said motif.
- 21. The composition of claim 20, wherein said oligonucleotide comprises a nucleotide sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, and SEQ ID NO:25.
 - 22. The composition of any preceding claim, further comprising one or more of the following proteins:
 - a protein disclosed in WO99/57280, or an immunogenic fragment thereof;

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- a protein disclosed in WO99/36544, or an immunogenic fragment thereof;
- a protein disclosed in WO99/24578, or an immunogenic fragment thereof;
- a protein disclosed in WO97/28273, or an immunogenic fragment thereof;
- a protein disclosed in WO96/29412, or an immunogenic fragment thereof;
- a protein disclosed in WO95/03413, or an immunogenic fragment thereof;
- a protein disclosed in WO99/31132, or an immunogenic fragment thereof;
- a protein disclosed in WO99/58683, or an immunogenic fragment thereof;
- a protein disclosed in WO99/55873, or an immunogenic fragment thereof; and/or
- a protein disclosed in GB-9928197.4, or an immunogenic fragment thereof.

10 23. A vaccine composition comprising:

- a) an immunostimulating amount of a Neisseria antigen; and
- b) an immunostimulating amount of an adjuvant composition comprising an oligonucleotide comprising at least one CG motif.
- 24. The vaccine composition of claim 23, wherein component (b) further comprises a second adjuvant.
 - 25. The vaccine composition of claim 23 or claim 24, wherein said peptide comprises SEQ ID NO:31.
 - 26. The vaccine composition of any one of claims 23 to 25, as further defined in any one of claims 2 to 21.

20 27. An adjuvant composition comprising:

- a) an oligonucleotide comprising at least one CG motif; and
- b) complete Freund's adjuvant.
- 28. The composition of claim 27 wherein said oligonucleotide comprises at least one phosphorothioate bond.
- 25 29. The composition of claim 27 wherein said oligonucleotide comprises a nucleotide sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

- SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, and SEQ ID NO:27.
- 30. The composition of claim 27 wherein said oligonucleotide comprises a CG motif flanked by two purines immediately 5' to said motif and two pyrimidines immediately 3' to said motif.
- 5 31. The composition of claim 30 wherein said oligonucleotide comprises a nucleotide sequence selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:12, SEQ ID NO:13, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:22, SEQ ID NO:23, SEQ ID NO:24, and SEQ ID NO:25.
- 32. A method of stimulating an immune response in a host animal comprising administering to said animal a composition of any one of claims 1 to 22 in an amount effective to induce an immune response.
 - 33. The method of claim 32 wherein said host animal is a mammal.
- 34. A method of immunizing a host animal against Neisseria infection comprising administering to said animal a composition of any one of claims 23 to 26 in an amount effective to induce a protective response.
 - 35. The method of claim 34 wherein said host animal is a mammal.
 - 36. The method of claim 35 wherein said mammal is a human.
- 37. A method of immunizing a host animal against *Neisseria meningitidis* comprising administering to said animal a composition of any one of claims 23 to 26 in an amount effective to induce a protective response, wherein said antigen is a Neisseria meningitidis group B peptide.
 - 38. The method of claim 37 wherein said peptide comprises SEQ ID NO:31.
 - 39. The method of claim 38 wherein said host animal is a human.
- 25 40. A composition as defined in any one of claims 1 to 29 for use a pharmaceutical.
 - 41. Use of a composition as defined in any one of claims 1 to 31 in the manufacture of a medicament for inducing an immune response in an animal.
 - 42. Use according to claim 41 wherein the composition is a composition as defined in claim 1 and the medicament is for inducing a protective immune response in an animal.